

Determination of the quality of a gas

Ins A17

Description

The invention concerns to a procedure for the photometric determination of the quality of gas, particularly of burnable gases, according to the pre-characterising part of claim 1 and claim 8 and also devices for the photometric determination of the quality of gas, particularly of burnable gases, according to the pre-characterising part of claim 12 and claim 23.

For the registration of the quality of gas for example in distribution networks for natural gas or the same already for a long time devices for the registration of the quality of the guided through gases are used, so-called devices for measuring the condition of gas. Natural gas shows because of being a natural product according to its origin and by mixture respective fluctuations in respect of its composition, whereas the composition for example of natural gas coming from the different hydrocarbons determines essentially the calorific value and there from extrapolated quantities. Therefore it is of great importance for the account of the guided-through amount of gas in a gas supply network and therewith the respective amount of energy, to determine exact the respective condition at the feeding point into the natural gas network and at the deliverance points of the customers and therewith to deduct a definite transported or supplied amount of energy. In doing so for the customer of the gas an invoice can always be stated according of the actual supplied amount of energy regarding to different conditions of the gas and a correspondingly varying amount of energy. Vice versa the detection of the condition of the gas offers the guarantee for the customer, to checkably obtain a desired quality and therewith a required amount of energy.

The registration of the quality of gas obtains additional relevance, since with the drop of the guiding-through monopoly the suppliers of natural gas guide through the same

SUB
A2
B3
09/869782
070501

network for delivering gases of quite different provenance and therefore also different composition. Only as far as possible easy and cost-effective detection of the condition of the gas by means of corresponding disposed cost-effective measuring devices and methods of measurement allows therefore a controllable and accurate accounting.

For the registration of the quality of gas as relevant quantities the standard volumetric gross calorific value $H_{v,n}$, the standard density ρ_n and the compressibility coefficient K have to be determined as correct as possible and also regarding the different gas qualities.

In practice for the energetic settlement of accounts first of all by means of flow measuring devices the transported volume V_b of the gas at working conditions (pressure p_b , temperature T_b) is measured. With knowledge of the condition of the gas the compressibility coefficient K can be determined, with which the volume of the gas V_n at standard conditions (pressure p_n , temperature T_n) is calculated.

$$V_n = \frac{p_b T_n}{p_n T_b} \frac{1}{K} V_b$$

By means of multiplication of this standard volume with the volumetric gross calorific value $H_{v,n}$ at standard conditions the transported amount of energy Q can be obtained:

$$Q = V_n H_{v,n}$$

Alternative the volume at working conditions V_b can directly be multiplied with the volumetric gross calorific $H_{v,b}$ at working conditions (Energymeter).

Another important quantity for applications with natural gas is the thermal output of gas burners; this varies in accordance to the gas quality and is characterised by means of the so-called Wobbe index W_v : gases with the same Wobbe index W_v deliver the same thermal output at a burner nozzle. For calculating the Wobbe index W_v the

standard density ρ_n of the gas is required, from which the relative density according to air is determined ($d_v = \rho_{\text{gas}} / \rho_{\text{air}}$)

$$W_v = \frac{H_v}{\sqrt{d_v}}$$

Therefore the determination of the gross calorific value $H_{v,n}$ at standard conditions has central relevance for the practical determination of the quality of gas for example for accounting purposes.

Until now different devices for the registration of the gas quality are used. So-called direct and so-called indirect procedures are known. By using direct procedures the quantities to be determined are measured separately and therefore the gas is transformed to standard conditions, by which partially expensive treatments of the gas are required.

At the most easiest way the gas condition can be determined by means of so-called calorimeters, in which by means of an open flame a gas probe is burnt and out the arising and to a cooling medium submitted heat quantity and the thereupon detectable temperature rise of a cooling medium the calorific value of the burnt gases can be determined. Such devices will need a complicated mechanic for the adjustment of a certain quantitative proportion of gas, combustion air and for example cooling air as cooling medium and are therefore expensive and error prone, especially relating to the open burning enhanced security requirements for the devices are necessary. Also the maintenance and calibration has to be carried out by qualified personnel, beyond that the calorimeter must be used in conditioned rooms. Therefore the purchase and operating costs of such test assemblies are very high.

Using calorimetry by means of catalytic burning (for example with pellistors) the probe gas is mixed with air and burnt at the 400 to 500°C hot helixes of a catalyst. The temperature rise of the catalyst is about proportional to calorific value. Because this procedure is based on a sensitive surface effect, it is subject strongish drifts and necessitates frequently calibration with search gas. The catalytic calorimeter are most

favorable of all here presented procedures, however they are better suited for control than for accounting because of their accuracy.

The direct measurement of the density ρ_b at working conditions is done in one way with hydrostatic balances, very expensive precision devices, with which the buoyancy of a ball filled with nitrogen is measured in accordance to the density of the surrounding medium, here of the probe gas. With another procedure a thin-walled metal cylinder, which is positioned by a current linkage of the probe gas, is set in oscillation. The density of the surrounding gas determines the resonant frequency of the cylinder, which is captured as sensitive measured quantity. Both procedures are very expensive for the determination of the standard density, because you they require an adjustment to the standard conditions.

The compressibility coefficient K can in this way not be measured directly, instead can be calculated by means of different numerical standard-arithmetic techniques out of the directly measurable gas quantities. One of these procedures, the so-called GERG88-procedure (DVGW-worksheet 486) needs thereby the input quantities listed in table 1. The amount of substance of CO₂ is determined according to the today's state of the art by a non dispersive infrared-spectroscopical procedure (NDIR), whereby the gas must be brought into a defined condition near or at standard conditions. The amount of substance of H₂ is practically of significance only when working with coke oven gases and can be practically left unattended in the typical natural gases today distributed in Europe. The compressibility coefficient K can be determined to 10⁻³ with the help of the GERG88-equation in case of sufficient accuracy of the input quantities.

P _b	pressure at working conditions
T _b	temperature at working conditions
ρ_n	density at standard conditions
H _{v,n}	volumetric gross calorific at standard conditions
x _{CO₂}	amount of substance of CO ₂
x _{H₂}	amount of substance of H ₂

table 1: input quantities of the GERG88-procedure

The other procedure for determination of the behaviour of real gases is done according to the AGA8-92DC-equation (ISO 12213-2:1997 (E)). This process requires as input quantities the amount of substance of 21 leading gas components (table 2) and gains just as an accuracy of 10^{-3} .

methane	CO ₂
ethane	N ₂
propane	H ₂ S
isobutane	He
n-butane	H ₂ O
isopentane	O ₂
n-pentane	Ar
n-hexane	H ₂
n-heptane	CO
n-oktane	pressure
n-nonane	temperature
n- decyl hydride	

table 2: input quantities of the AGA8-92DC-equation

The state of the technology includes besides the direct measurement techniques also the indirect measurement of the gas quality by means of gas chromatography. Thereby a defined volume of the probe gas is brought into a defined condition and is carried by a carrier gas, typically helium, through a system of gaschromatographic separation columns. On account of their different times of retention the individual gas components reach the downstream sensor, generally this is a detector for caloric conductivity, at the end of the separation column separated in time. The peak area of the sensor signal can therefore be interpreted as amount of substance, whereas the evaluation must be carried out in comparison with a reference gas, that must have just about a similar composition like the probe gas. The drawback of the gas chromatography has its reason in the expensive sample preparation and installation of the whole system, and in the expensive maintenance and operation by well-trained personnel. From the amounts of substances of the individual gas components, as the gas chromatography delivers, all relevant gas quantities can be calculated. For the implementation of such indirect measurements in the way of chromatography automatically working process

chromatographs with detectors for caloric conductivity are deployed. These devices measure normally eleven components of the natural gas (N_2 , CO_2 , CH_4 , C_2H_6 , C_3H_8 , C_4H_{10} , C_5H_{12} , C_6+ and so on). As carrier gas helium will be used, whereby its light volatility in practice often leads to prematurely emptying of the bottle for the carrier gas and therefore leads to short maintenance cycles for such a device for measuring the quality of a gas. As calibrating gas a gas is chosen, that is similar to the natural gas to measure. Such chromatographic systems carry out measurement cycles without interruption, in order for capturing changes in the quality of the gas immediately. These leads to a high consumption of carrier gas and calibrating gas and has moreover the effect, that maintenances of the device have to be performed in relatively short intervals.

It is also known, to determine the composition of a gas with conventional infra-red-gas-analyzers. Such analysers working in the middle infra-red or near infra-red area offer however not the requested requirements to high precision and in particular to stability for a determination of the caloric value under the measurement conditions which are required here. Also always a parallel reference measurement is to be carried out beside the intrinsic measurement of the probe gas, for the sake of compensating the at least essential influences of failures. As a measuring result the known infra-red-gas analysers deliver superimposed frequency spectrums, that make very difficult a conclusion to individual components of a inspected gases, if not even forbid such a conclusion.

In the literature also has been described a infrared spectroscopical procedure for gas analysis („Optical BTU sensor Development“, Gas Research Institute GRI-93/0083), that determines by means of so-called multivariate analysis (MVA) of the near infrared spectrum of gases the volumetric concentration of the amount of substances of the containing carbon components of the gas and therewith of the volumetric calorific value under operating conditions. This procedure delivers however not the calorific value $H_{v,n}$ under standard conditions and not the standard reference density ρ_n and the amount of substance of CO_2 , so that it is not qualified for the determination of the

compressibility coefficient K and therefore not for the complete determination of the quality of a gas. By the determination of the calorific value by means of known photometric methods moreover minor claim to the necessary instrumental equipment for the realisation of the procedures are required, whereby a benefit in the speed of the coverage of the absorption spectrums of the natural gas in near or middle infrared spectral area is obtained. The entire absorption spectrum of the natural gas is therefore put additive together from of the sum of single spectrums of components represented in gas and therefore can be measured and be analysed with the aid of more appropriate methods of spectral analysis. Doing this the ascertained quota of extinction of a component in the entire spectrum of the natural gas is in effect equivalent to the part of the concentration of this component within search gas (so-called Beer-Lambert-law). With the knowledge of the calorific value of this respective component the calorific value of the entire mixture of gases can then be calculated as a summation value. Problematically in this procedure of spectral analysis is however the intense overlap by absorption bands of different components, which frequently lead to inaccurate results and beyond that to a high calculation effort.

One more infrared spectroscopical procedure concerning to the DE 198 38 301 A1 will act as a direct spectral evaluation (DSA) with a spectral function, with which the spectrum of the gas is folded. The procedure allows the specification of the volumetric calorific value $H_{v,b}$ under operating conditions direct from the spectrum. Herewith it is utilized, that while burning the gas the respectively caused heat of the reaction is based on the combustion of C-H-bindings and a thereby caused heat quantity depends to the present binding energy. This is thereby exploited, that the oscillations of the C-H-bindings, which show an equal to each other, certain binding energy and produce the same heat quantity during a combustion, interact with an associated wavelength of an electromagnetic radiation. Hereby it is possible by means of a wavelengths-resolved measuring and a wavelengths-dependent valence of the grade of interaction of these oscillations to calculate the calorific value $H_{v,b}$ of the gas, without the requirement of an identification of individual gas components. In this document it is additionally

proposed a device for the realisation of such a procedure, which is tuned for the specific requirements of the measuring method and enables a weighted summing up of the grades of interaction. With this procedure indeed the calorific value $H_{v,b}$ of a mixture of gases under operating conditions can be determined, the other quantities relating to the determination of gas quality can itself not be determined with measuring techniques.

After having today's state of the art there is no, especially no uniform procedure, that determines under operating conditions the substantial quantities volumetric standard-calorific value $H_{v,n}$, standard-density ρ_n and compressibility factor K .

Therefore it is object of the pending invention, to propose procedures and devices for the determination of the quality of gas, with which the substantial quantities volumetric standard-calorific value $H_{v,n}$, standard density ρ_n and compressibility factor K can be determined by means of a spectroscopical registration of the quality of the gas as well as respective evaluations.

The solution of the object according to the invention results in respect of the procedures from the characterising features of the claims 1 respectively 8 in accordance with the features of the associated pre-characterising part. The solution of the object according to the invention results in respect of the devices from the characterising features of the claims 12 respectively 23 in accordance with the features of the associated pre-characterising part. Further advantageous embodiments of the invention result from the respective dependent claims.

The invention concerns to a first procedure according to claim 1, with which the determination of the quality of gas of a probe gas, in particular a burnable gas, is carried out based on a spectrum of the probe gas determined under operating conditions by means of infrared spectroscopical measurement procedures. Herein the quality of gas of a probe gas is determined in such a way according to the invention, that in a first step of the procedure the amounts of substances x_i of the components of the probe

00000000000000000000000000000000

gas at operating conditions are determined out of the spectrum, where after default values for compressibility factor K and real gas factor Z_n are preset for calculation of the wanted compressibility factor K and than out of quantities at operating conditions of the probe gas as well as from the amounts of substances x_i and substance specific quantities such as calorific values per sort of molecule of the components and the respective masses of molecules, and taking into account of the selected default values for compressibility factor K and real gas factor Z_n the needed input quantities for the determination of the compressibility factor K are determined. These input quantities are used, to calculate the compressibility factor K by means of standard-arithmetic procedures. In a further step of the procedure an iterative calculation in the way of an iterative recalculation of the input quantities is carried out with the determined value for the compressibility factor K as long, until the value of the compressibility factor K converges. There from the volumetric standard calorific value $H_{v,n}$ and the standard density ρ_n can be calculated. In the case of converging the needed quantities compressibility factor K , standard calorific value $H_{v,n}$ and the standard density ρ_n for determination of the quality of gas of a probe gas are known and can be used respectively for example for the calculation of the energy content of the transported gas. At each iteration the just determined value for the compressibility factor K is put back again into the equations for calculating the input quantities of the standard-arithmetic procedure and a new iteration step is carried out. By means of this procedure starting with the determination of the amounts of substances x_i out of the recorded spectrum and the iteration procedure with the help of the default values for the compressibility factor K and real gas factor Z_n after a respective number of iteration steps the really existing values for the compressibility factor K and there from than the values for standard-calorific value $H_{v,n}$ and the standard density ρ_n in the probe gas can be determined with adequate accuracy. In this way the compressibility factor K is determined out of the spectroscopical measurable quantities in operation conditions as well as out of the default values for the compressibility factor K and real gas factor Z_n and the real gas calculation itself can be carried out with standard-arithmetic procedures and these arithmetic procedures itself are again iterational procedures.

Therefore there is a base of a two-stage iteration, in which in a first iterative step with the default values for the compressibility factor K and real gas factor Z_n and the input quantities of the standard-arithmetic procedure the compressibility factor K is calculated by means of iterative procedures and in the case of non-convergence the compressibility factor K and the quantities, which can be determined again there from, are put back after a new calculation of the input quantities of the standard-arithmetic procedure as input quantities for a new iterative loop. Beneath the convergence of the compressibility factor K the convergence of the standard-calorific value $H_{v,n}$ and the standard density ρ_n can be proofed. This procedure owns the advantage, that out of the quantities being measurable at operating conditions and the spectrum, which can be determined direct at the operating conditions, the standard values of the probe gas can be determined directly without subjecting the probe gas to the otherwise necessary, expensive treatments for transferring it into standard conditions.

In an especially preferred embodiment as a standard-arithmetic procedure for the determination of method of iteration AGA8-92DC is used, which is committed as an international uniform arithmetic procedure in ISO 12213/2. As significant input quantity for this procedure AGA8-92DC here the molar amounts of substances of the infrared active components and the molar amount of substance of nitrogen N_2 is used.

In an other preferred embodiment as a standard-arithmetic procedure for the determination of the compressibility factor K the method of iteration GERG88 is used, which is committed in the DVGW-working sheet 486 and described detailed. This method of iteration allows also the determination of the compressibility factor K , wherein as input quantities out of the molar amounts of substances of the infrared active components the standard-calorific value $H_{v,n}$ and the standard density ρ_n and the concentration of CO_2 can be calculated.

In an especially preferred embodiment the determination of the amounts of substances x_i of the infrared active components of the probe gas at operating conditions is carried out starting from the recorded spectrum by means of a procedure according to the multivariate analysis (MVA), which is described in detail in the report of the Gas

Research Institute GRI-93/0083 „Optical BTU Sensor Development“. By using this procedure of multivariate analysis (MVA) out of the spectrums of the probe gas, recorded at operating conditions, the amounts of substances of the infrared active components are determined and are therefore available for the calculation of the compressibility factor K.

In a preferably embodiment the default values of the compressibility factor K and the real gas factor Z_n are taken from a characteristic diagram, that describes the influence of the pressure p_b at operating conditions and the temperature T_b at operating conditions for a known composition of a gas similar to the composition of the probe gas. By doing this the fact is used, that the pressure p_b at operating conditions has a great influence, the temperature T_b at operating conditions as also the real composition of the gas, especially for usual compositions of burnable gases, have a minor influence at the values of the real gas factor Z_n . Therefore it is possible with characteristic diagrams of a gas similar to the composition of the probe gas, which have been determined before and in which the relation between pressure and temperature is outlined, to take a starting value for the iterative calculation according to a standard-arithmetic procedure as a good first approximation for the pressure p_b at operating conditions and the temperature T_b at operating conditions, which converges with a real good accuracy after a small amount of iteration steps to the real compressibility factor K.

Furthermore it is especially preferable, that directly from the recorded spectrum the amounts of substances of the single burnable components of the probe gas at operating conditions and the amount of nitrogen N_2 of the probe gas are determined as a function of the amounts of substances of the infrared active components of the probe gas. Therefore as an additional information the pressure of the probe gas, in which the amount of substance of nitrogen is also taken into consideration, has relevance for the calculation.

In further embodiment the amount of substance of nitrogen N_2 and the amounts of substances of the infrared active components complements each other resulting in the total volume of the probe gas, in which further substances, as for example oxygen O_2 , hydrogen H_2 and water for example are contained in typical burnable gases only in small traces and therefore can as a rule be neglected.

The invention concerns furthermore to a procedure for the determination of the quality of gas of a probe gas, in particular a burnable gas, proceeding from a spectrum of the probe gas determined under operating conditions by means of infrared spectroscopical measurement procedures.

This procedure will hereby be developed further in a way according to the invention, that in a first step of the procedure default values for compressibility factor K and real gas factor Z_n are preset for calculation of the wanted compressibility factor K , in a further step of the procedure input quantities for the determination of the compressibility factor K are determined by using the pressure p_b at operating conditions and the temperature T_b at operating conditions of the probe gas, which can be determined without problems from the operating condition of the probe gas, and out of the values for the calorific value $H_{v,b}$ at operating conditions and the density ρ_b at operating conditions, whereby as further input quantity the molar amount of substance of CO_2 is determined by means of a further absorption band of the spectrum.

Hereby both the calorific value $H_{v,b}$ at operating conditions as also the density ρ_b at operating conditions can be directly determined based on the spectrum, in which simultaneously based on the same spectrum the further absorption band for the amount of substance of CO_2 can be determined. With these input quantities in a further step of the procedure the compressibility factor K is calculated by means of the iterative procedure GERG88. On the base of theses results of the iterative procedure GERG88 than it is examined, if the value for the compressibility factor K converges and than there from the volumetric standard calorific value $H_{v,n}$, the standard density ρ_n and the amount of substance of CO_2 is calculated. If a convergence is

ascertainable, than as results of the procedure according to the invention the determined compressibility factor K as also the therewith calculated values for the standard calorific value $H_{v,n}$, the standard density ρ_n and the amount of substance of CO_2 can be used directly for further purposes. If a convergence is not ascertainable, than an

5 iterative recalculation of the input quantities is carried out with the determined value for the compressibility factor K as long, until the convergence of the compressibility factor K and if necessary the standard calorific value $H_{v,n}$ and the standard density ρ_n is to be noticed. During each iteration the just determined value of the compressibility

10 factor K is put again into the equations for the calculation of the input quantities of the standard-arithmetic procedure and a new iteration step is carried out. The procedure according to the invention shows the advantage, that also without knowledge of the exact amounts of substances of the single components contained in a probe gas the wanted target quantities at working conditions can be solely determined out of the spectroscopical quantities at working conditions and the start values for the

15 compressibility factor K and the real gas factor Z_n , in which a two-staged iterational approach, on one hand the calculation by iteration of the compressibility factor K while carrying out the procedure according GERG88, on the other hand with an iteration while recalculating again the input quantities for GERG88 on base of the in the step before determined compressibility factor K until the convergence of the results for the

20 compressibility factor K and the volumetric standard calorific value $H_{v,n}$ and the standard density ρ_n can be carried out. This so-called direct spectral evaluation (DSA) is fundamentally known in the state of the art for the determination of the calorific value $H_{v,b}$, in which the calculation of the density ρ_b at working conditions and the compressibility factor K solely on the base of the recorded spectrum was not known

25 until now.

It is most preferable, that the calorific value $H_{v,b}$ at operating conditions and the density ρ_b at operating conditions can be determined by means of spectral functions for weighting of a value directly from the spectrum of the probe gas, in which for this it

30 can be used, that based on the oscillation frequency of the binding of the infrared active

components of the gas their spectral position can be seen and this again relates to the reduced mass of the binding partners. With this the spectrum contains in its amount and its spectral distribution also information for the determination of the density ρ_b at working conditions, if it is suitably evaluated.

5

In a further embodiment with the spectral functions for weighting of a value the weighted influence of the amounts of substances of the components of the probe gas is described for the calorific value $H_{v,b}$ at operating conditions and the density ρ_b at operating conditions. In this case the single amounts of substances have not to be determined explicitly because of the spectral functions for weighting of a value, but the influence of these amounts of substances is represented in the spectral functions for weighting of a value.

10

Also it may be thinkable, that the default values for compressibility factor K and real gas factor Z_n are taken from a characteristic diagram, that describes the influence of the pressure p_b at operating conditions and the temperature T_b at operating conditions for a known composition of a gas similar to the composition of the probe gas. As described before for the procedure according to claim 1, in this way good starting values as a good first approximation for the compressibility factor K and the real gas factor Z_n can be determined, which contributes to a fast convergence also of this procedure according to the invention.

20

The invention also concerns to a photometric device for the determination of a transmission spectrum of a probe gas, especially for carrying out one of the procedures according to claim 1 or claim 8.

25

The invention bases on a radiation source emitting a measurement radiation, in which the measurement radiation passes through a probe cell for capturing a probe gas and enters after passing through a modulation unit for modulating the measurement radiation into at least one radiation receiver, which generates electrical measurement signals according to the incoming intensity of the measurement radiation and transmits

30

SUB
AS
13
TUM

these to an electronical unit, which determines a transmission spectrum out of the measurement signals, wherein the modulation unit shows a spectral switch unit. In a way according to the invention, the radiation source emitting a measurement radiation is connected with the probe cell and the probe cell with the spectral switch unit by means of at least each one light guiding device. By this way a guiding of the measurement radiation can be obtained, which is free of free air gaps in the device respectively near to the probe cell and in these air gaps no interference influence can affect the guiding.

Also between on the one hand the radiation source emitting a measurement radiation and the spectral switch unit as well as the probe cell a three-dimensional separation can be provided, which can be bridged by means of the light guiding device. By this it is obtained, that the real measurement device can be placed apart from the coupling to the for example pipeline guiding the probe gas, which can be not easily to be accessible or be subjected to contaminations or the same harmful influences. Also the technical realisation of the device can be further simplified.

In a further embodiment it may be thinkable, that in the area of the three-dimensional separation between on the one hand the radiation source emitting a measurement radiation and the spectral switch unit as well as the probe cell an explosion-proofed barrier is provided. This may be of importance in areas, in which a special necessity for a fire barrier or explosion proofness exists.

A further embodiment shows, that the spectral switch unit shows optical filters, with which regions of the transmission spectrum are extracted by filtering and are guided to the one or the respective radiation receivers. In this way a well-aimed evaluation of the spectrum with less instrumental effort and low requirements for the evaluation of the determined data is possible. Particularly simply is the evaluation, if in a further development the optical filters and their filter regions are chosen in such a way, that they record directly spectral regions for evaluation with a procedure of the direct

spectral evaluation (DSA). For example the spectral switch unit could be provided with each a filter at least for CH₄, higher CH-compounds, CO₂ and a reference gas.

In a preferable way for each region of the spectrum to be filtered a special optical filter and a respective radiation receiver can be provided. Also it may be thinkable, that for

5 each region of the spectrum to be filtered a special optical filter is provided, which is arranged in a filter carrier, which is relatively movable relative to the radiation receiver. In this way the costs for expensive radiation receiver can be further reduced.

10 It is a special advantage for the operational reliability of the device, if for the adjustment of the optical system of the device a spectroscopical inactive inert gas, preferably nitrogen N₂, is passable into the probe cell. Also for the examination of the measured values of the device a defined calibration gas is passable into the probe cell. By means of both of these two measures the long-time operation of the device according to the invention can be assured better without the possibility, that the long-
15 time changes at the device may result in erroneous measurement results. Therefore for example the measurements for adjustment of the optical system and/or calibration of the measured values can be carried out in definable time intervals.

20 The invention concerns furthermore to a generic photometric device for the determination of a transmission spectrum of a probe gas, especially for carrying out one of the procedures according to claim 1 or claim 8.

25 Such a generic device can be developed further, if the spectral switch unit shows a chopper arrangement, which transmits because of their selective transmission behaviour only specific spectral regions of the spectrum in the measurement radiation caused by the probe gas to the radiation receiver. In this way in further development the chopper arrangement can be provided with such a transmission behaviour, that the transmitted spectral regions are suitable for the further evaluation by procedures of the direct spectral evaluation (DSA). By means of the chopper arrangement only that part
30 of the whole spectrum is measured, which is of interest for the determination of the

09869732-070501

SUB 7
ALP 20

quality of gas for example of a probe gas of typical composition at all und beneath with simple mechanical means allows an exact selection of these spectral regions. Also such a chopper arrangement is build up easily and mechanically solid and insensitive.

5 In a first embodiment chopper arrangement is provided with a rotating aperture with free sector elements, which release regions of the measurement radiation in respect of the radiation receiver, which correspond to appointable spectral regions of the measurement radiation. By this the respective allocation of the sector elements is fixed to the spectral regions, which shall be recorded sequentially by the radiation receiver while the rotation of the aperture and at which the release of the regions of the measurement radiation is carried out sequentially for single spectral regions.

10 In an alternate embodiment the chopper arrangement can be provided with an aperture with a spiral opening, in which the release of the regions of the wavelength of the measurement radiation is caused continuously for the whole spectrum. By this means also compositions of probe gases can be determined, whose spectrums have to be captured over wide regions, also a higher flexibility for the evaluation of the spectrums can be obtained by this.

20 There is an especial advantage, if the released wavelength of the measurement radiation, which passes through the chopper arrangement, can be obtained by means of capturing the rotational position of the aperture. This can be realised in a simple manner for example by a shaft encoder, which is build up mechanically simple, inexpensive and robust and allows a secure synchronisation of the capturing of the spectrums with the respective wavelength.

25 In a further development the chopper arrangement is provided with two groups of sector elements alternatively releasing the measurement radiation, in which a first optical waveguide guides the measurement radiation released by the sector elements of the first sector element group into the probe cell and after passing through the probe cell to the radiation receiver and a second optical waveguide guides the measurement

30

radiation released by the sector elements of the second sector element group directly to the radiation receiver. At the here especially based application very high requirements are to be fulfilled in respect of the long-time stability of the device against signal drift. The following three factors contributes to a signal drift: fluctuation in respect of the luminous power of the radiation source, fluctuation of the sensitivity of the radiation receiver (both for example as result of aging processes) and fluctuation of the CO₂-concentration in the surrounding air, which results in an instable CO₂-underground spectrum. The CO₂-concentration in the surrounding air and therewith in the free beam path of a device varies for example on behalf of the presence of persons in the surrounding. All three influences can be compensated by means of the here proposed dual-beam building up of the device, with which a permanent comparison of the signal with the reference signal can be carried out. By means of the sector elements of the sector element groups the each released measurement radiation of that waveguide can be used, which is guided to the radiation receiver, as reference for eliminating the influence of existing CO₂ in the surrounding of the probe cell and/or the device that changes the radiation source and/or the radiation receiver. Beneath the elimination of the influence of CO₂ in the surrounding in this way also changes of the radiation source and/or the radiation receiver can be eliminated, which can be the result of aging processes or contaminations.

In this way in a first embodiment the measurement radiation released by the sector elements of the sector element groups can be concentrated by means of first and second optical waveguide into one or more filters or a dispersive element, preferably a monochromator.

There is a special advantage, if the radiation receiver collects the measurement radiation, which is coming out of the one or more filters or the dispersive element and each released through the sector elements of the sector element groups of both optical waveguides.

In a further embodiment it may be possible, that the measurement radiation, which is each released through the sector elements, is guided through the first and the second waveguide to the input of the one or more filters or the dispersive element, in which at the chopper arrangement also available further sector element groups lock on the measurement radiation, which is released of the one or more filters or the dispersive element, alternatively to the radiation receiver. By theses means a reliable synchronisation of the modulations can be assured using a mechanically simple build up arrangement of the sector element groups of the aperture. In a further variant it may be thinkable, that the chopper arrangement carries out both the selection of the wavelengths for the spectrum as well as the alternating reverse of the measured section between the waveguides.

In this case a further development may be possible, in that the measurement radiation, which is each released through the sector elements, is guided together by means of the first and the second waveguide in a Y-fibre coupler, which guides the measurement radiation of the first and the second waveguide to the one or more filters or the dispersive element. This allows an especially simple design of the filter and the dispersive element and the concentration of the splitted beams.

In further development it may also be thinkable, that the probe cell is sweepable with an infrared inactive gas, preferably nitrogen N_2 , for carrying out a null measurement for the compensation of dirt accumulation or the same of the optical facilities of the device. In this way it is possible to capture with this probe cell filled with gas of definite and known properties, if there are deviations relative to an initial state and these deviations can be taken into consideration for further measurements.

A preferred embodiment of the invention provides, that the probe cell contains an optical hollow shaft guide, into which the probe gas can be lead. In this case as advantage can be cited, that the probe cell can be coupled as hollow shaft guide very simple and without additional optics to the supplying and the carrying away

waveguides. The build up is much more compact as in the case of a free guided parallel beam. There is as further advantage, that the free effective length of path of the measurement radiation through the probe cell at passing through a probe cell provided with a hollow shaft guide is enlarged related to conventional probe cells because of multiple reflections at the vaporized inner walls of the hollow shaft guide.

Especially for the mounting of such hollow shaft guides there is as advantage, that the cross-sectional dimensions of the hollow shaft guide are adjustable in general the same or less larger as the cross-sectional dimensions of the optical waveguide, for being able to be coupled without additional optics. Also the hollow shaft guide can be coupled directly with the optical waveguide.

The invention provides also, that for the coupling in of the measurement radiation into a probe cell the probe cell is providable with lenses with a radial gradient of refraction index, so-called GRIND-lenses. In the atmospheric air gases as CO₂ and water steam are contained, which can cause trouble because of their fluctuation the gas measurement in the probe cell. Therefore it has to be avoided, that the measurement radiation is guided through free optical paths with surrounding air, whereas in a conventional probe cell also free distances cannot be avoided in the fibre optic, which can be avoided by using GRIND-lenses.

A further preferable embodiment of the invention provides, that the measurement radiation used in the photometric device can be modulated according to amplitude and/or wavelength. Herewith in further development the radiation source itself shows devices for modulating the measurement radiation, also it is possible, that the measurement radiation passes through an additional modulation unit after leaving the radiation source, which than modulates the measurement radiation. Such a spectral modulation is known for the expert and can be realized by means of optical gratings, an optical prism, optical filters, prisms or gratings in combination with chopper units or the same, in which it is of specific advantage, if the modulation is carried out in such a way, that as starting quantities the direct spectral regions result, which are usable for

the procedures of direct spectral evaluation (DSA). In relation to this well-aimed choice of specific spectral regions is referred to the above described preferable embodiments. An especially preferred embodiment of the procedures according to the invention and the device according to the invention shows the drawing.

5

It is shown in

- Sub 7**
A7
B7
09869782-070501
- 10
- Figure 1 - a typical distribution in a spectrum for different ingredients of natural gas,
- Figure 2a - fundamental procedure of a two-staged iterational procedure according to claim 1,
- Figure 2b - fundamental procedure of a two-staged iterational procedure according to claim 8,
- Figure 3 - a fundamental structure of a photometric device according to the state of the art,
- 15 Figure 4 - a device according to the invention related to claim 12, which is coupled to a probe cell by means of waveguides and determines the spectrums by means of filters,
- Figure 5 - a first device according to the invention related to claim 23, in which a chopper arrangement with a sector element aperture is provided for selection of wavelengths,
- 20 Figure 6 - a second device according to the invention related to claim 23, in which a chopper arrangement with a spiral aperture is provided for selection of wavelengths,
- Figure 7 - a further device according to the invention related to claim 23 according to the reference beam principal of claim 29,
- 25 Figure 8 - an embodiment of the device according to figure 7 with synchronisation by the chopper arrangement itself,
- Figure 9 - an embodiment of the probe cell as hollow shaft guide,
- Figure 10 - a connection of the probe cell to a waveguide with the help of GRIND-lenses,
- 30

Sub 11
CS 20
Figure 11 - an embodiment of the invention according to claims 41 to 43 with a modulation of the measurement radiation.

In figure 1 it is described, which distribution can be found in the spectrum for different components contained in natural gas.

Natural gas as distributed typically in Europe consists generally of the components as methane, ethane, propane, butane, pentane and higher hydrocarbons such as carbon dioxide and nitrogen. According to DIN 51857 mixtures of gases count as similar to natural gas, if their amounts of substances fulfil the following conditions:

$x(\text{CH}_4) \geq 0.5$
$x(\text{N}_2) \leq 0.3$
$x(\text{CO}_2) \leq 0.15$
$x(\text{C}_2\text{H}_6) \leq 0.15$
$\sum x_i \text{ over all other components} \leq 0.05$

Table 3: limiting value for gases similar to natural gas according to DIN 51857

The remaining components are mostly higher hydrocarbons with amounts of substances of about 10^{-3} , further substances as O_2 , H_2 , water and other occur in typical natural gases in trace amounts, which do not disturb a measurement with the demanded accuracy. With exception to nitrogen all relevant components of mixtures of gases similar to natural gas are infrared active, so that they absorb radiation in the infrared spectral region and can be detected in this way.

The extinction coefficients of the relevant gas components are described in figure 1, while the components methane and CO_2 are isolated in specific spectral regions, the higher hydrocarbons overlap spectral very much. The figure 1 shows moreover typical spectral regions for the determination of the single components (CH_4 , CO_2) respectively groups of components (C_2H_6 and higher hydrocarbons).

According to the Beer-Lambert-law the infrared components contribute to the extinction in the infrared spectrum according to their particle number density in the optical measuring volume. Based on the spectrum the volumetric amounts of substances of the infrared substances of an unknown gas probe can be determined at measuring respective working conditions.

For determining the volumetric calorific value at working conditions these information are sufficient, because the „invisible“ nitrogen does not contribute to the calorific value. According to the state of the art for this two procedures are described. According to the one procedure (DVGW-working sheet 486) the analyses of the gas mixture is carried out referring to their composition of single components. Therefore the amounts of substances of the single burnable components are determined out of the spectrums by means of the mathematical procedure of multivariate analyses (MVA) and the calorific value is calculated as corresponding weighted sum. In the second procedure (direct spectral analyses DSA) (ISO 12213-2:1997(E)) not single components are identified, but by means of a convolution integral with a specific spectral function the calorific value at working conditions is calculated directly out of the spectrum.

In practice the measurement of the quality of gas not only the calorific value $H_{v,b}$ at operation conditions is expected, but the calorific value $H_{v,n}$ at standard conditions and the standard density ρ_n and the compressibility factor K . For the calculation of these quantities the knowledge about the amount of substance of nitrogen in the gas is required, but this cannot be gained directly out of the infrared spectrum.

The procedures introduced here make possible a determination of the relevant gas quantities out of the infrared spectrum without explicit knowledge of the amount of substance of nitrogen. Thereby the contribution of nitrogen is printed out as function of the other spectroscopical measurable components; as additional information supplementary the pressure of the gas enters the figure, which also takes into account the amount of substance of nitrogen. Basis for this is the gas equation of the real gas

with the pressure p , the volume V , the real gas factor Z as function of the condition (p, T) and the composition of gas x_i , of the sum of all the molecules N , the Boltzmann factor k_B and of the temperature T .

$$p V = Z(p, T, x_i) N k_B T$$

With this the mathematical representations of the wanted quantities for a first procedure according to claim 1 for the determination of the compressibility factor K result in the following equations. These representations contain however the wanted compressibility coefficient K as well as the unknown real gas factor $Z_n = Z(p_n, T_n)$ of the unknown gas at standard conditions.

The calculation is achieved by one in Fig 2 nearer explained two-stage iterational approach, in which the appropriate starting values for K and Z_n are selected. For the determination of the compressibility factor K either the procedure GERG88 (DVGW-worksheet 486) or the procedure AGA-92DC (ISO 12213-2:1997(E)) will be used. The respective input quantities of the two iterational procedures are calculated out of the spectrum analysis of the gases with the applicable selected starting values for Z_n and K .

The input quantities for the standard-arithmetic procedure AGA8-92DC are described with the volumetric concentrations of the amounts of substances of the infrared active components and the starting values for K and Z_n as following:

$$x_i = \frac{k_b T_b}{p_b V_n} K Z_n N_i$$

$$x_{N_2} = 1 - \frac{k_b T_b}{p_b V_b} K Z_n \sum_i N_i$$

Herein the x_i indicate the amounts of substances of the infrared active components and x_{N_2} the amount of substance of nitrogen.

The input quantities for the alternative standard-arithmetic procedure GERG88 are calculated with the volumetric concentrations of the amounts of substances of the infrared active components and are described with the starting values for K and Z_n as follows:

$$H_{v,n} = \frac{p_n T_b}{p_b T_n} K \sum_i H_{m,i} N_i$$

$$\rho_n = \frac{p_n T_b}{V_b p_b T_n} K \sum (m_i - m_{N_2}) N_i + \frac{p_n m_{N_2}}{T_n k_B Z_n}$$

$$x_{CO_2} = \frac{k_B T_b}{p_b V_b} K Z_n N_{CO_2}$$

with calorific values $H_{m,i}$ per molecule of the component i and the respective masses of molecules m_i , the mass of the molecule of nitrogen m_{N_2} and the number of CO_2 -molecules N_{CO_2} in the measurement volume.

A multivariate analysis (MVA), as outlined in the literature (DVGW-worksheet 486), delivers the volumetric amounts of substances of the gas components at operation conditions. The here presented procedure according to claim 1 describes, how moreover with these data the standard calorific value $H_{v,n}$, the standard density ρ_n and the compressibility factor K can be ascertained.

With the procedure according to claim 8 by means of a spectral weighting function the spectrums can be evaluated directly without detailed resolution of the individual gas components. The specification of the calorific value $H_{v,n}$ at operation conditions according to this procedure is lined out in the patent application DE 198 38 301.0. The

volumetric amount of substance of CO₂ at operating conditions can be evaluated as separate absorption band according to common spectroscopic procedures.

Vital component of the here introduced procedure according to claim 8 is the spectral measurement of the density ρ_b at working conditions, therefore also the convolution of the spectrum with a spectral function is used. The physical background of the procedure is herein, that each binding of the infrared active gas components contribute for extinction and so represents the mass of the partners of the binding in the spectrum. The frequency of oscillation of the binding and therewith their spectral location depends on the reduced mass of the partners of the binding. Therewith the spectrum contains in its amount and its spectral distribution at appropriate evaluation information for the determination of the density of the gas.

The input quantities for the herein used standard-arithmetic procedure GERG88 result thereby out of the direct evaluation of the spectrums and with the starting values for K und Z_n:

$$H_{v,n} = \frac{\rho_n T_b}{\rho_b T_n} K H_{v,b}$$

$$\rho_{v,n} = \frac{\rho_n T_b}{\rho_b T_n} K \rho_{v,b}$$

$$x_{CO_2} = \frac{k_B T_b}{\rho_b V_b} K Z_n N_{CO_2}$$

In the figures 2a and 2b the flows of the two-stage iterational procedure according to the above mentioned equations are illustrated once more precisely. In the figure 2a is represented the principal flow of the calculation of the procedure according to claim 1 with the both alternative iterational procedures AGA8-92DC and GERG88, which start from the results of the multivariate analysis (MVA) by means of the ascertained

Sub 1A9
C1217

spectrum. In the figure 2b is represented however the principal flow of the calculation of the procedure according to claim 8, that starts from the results of the direct spectral evaluation (DAS) and than uses the iterational procedure GERG88.

5 At the procedures according to figure 2a a first approach for the values of K and Z_n is made, according to the determination of the directly with conventional means measurable quantities of the probe gas as pressure p_b and temperature T_b at operating conditions as well as of the evaluation of the spectrum by means of the multivariate analysis (MVA), which at most easy can be drawn from a characteristic diagram, that
10 represents the influence of pressure p_b at operating conditions and temperature T_b at operation condition for a known composition of gas similar to the probe gas. With these quantities then for the first time according to a chosen standard-arithmetic procedure (AGA8-92DC or GERG88) the respective input quantities of the respective standard-calculation procedure will be calculated and these are then used in the
15 respective iterational procedure of the standard-arithmetic procedure. As a result of this a value for the compressibility factor K is obtained as well as the here from calculatable quantities standard calorific value $H_{v,n}$ and standard density ρ_n . Converges the compressibility factor K, and if necessary also the standard calorific value $H_{v,n}$ and standard density ρ_n , towards a final value, than the calculation of the wanted quantities
20 is completed, because these are now directly available respectively can itself be calculated by means of the calculated values for the compressibility factor K. Is there however no convergence to be ascertained, than the result of the compressibility factor K is put back again into the input quantities of the standard-arithmetic procedure and with these changed input quantities the iteration is started once again and after passing
25 through the steps of the procedure once again examined for convergence. Doing this beside the convergence of the value of the compressibility factor K, like this is represented in the figures 2a and 2b, additionally also the convergence of the quantities evaluable from the compressibility coefficient K, for example standard calorific value $H_{v,n}$ and standard density ρ_n . can be examined and be utilised as further criterion for
30 termination of the iteration.

At the procedure according to figure 2b a first approach for the values of K and Z_n is made, according to the determination of the directly with conventional means measurable quantities of the probe gas as pressure p_b and temperature T_b at operating conditions as well as of the evaluation of the spectrum by means of the direct spectral evaluation (DAS), which at most easy can be drawn from a characteristic diagram, that represents the influence of pressure p_b at operating conditions and temperature T_b at operation condition for a known composition of gas similar to the probe gas. With these quantities then for the first time according to the standard-arithmetic procedure GERG88 the respective input quantities of the respective standard-calculation procedure will be calculated and these are then used in the respective iterational procedure. As a result of this a value for the compressibility factor K is obtained as well as the here from calculatable quantities standard calorific value $H_{v,n}$ and standard density ρ_n . Converges the compressibility factor K , and if necessary also the standard calorific value $H_{v,n}$ and standard density ρ_n , towards a final value, than the calculation of the wanted quantities is completed, because these are now directly available respectively can itself be calculated by means of the calculated values for the compressibility factor K . Is there however no convergence to be ascertained, than the result of the compressibility factor K is put back again into the input quantities of the standard-arithmetic procedure GERG88 and with these changed input quantities the iteration is started once again and after passing through the steps of the procedure once again examined for convergence. Doing this beside the convergence of the value of the compressibility factor K , like this is represented in the figures 2a and 2b, additionally also the convergence of the quantities evaluable from the compressibility coefficient K , for example standard calorific value $H_{v,n}$ and standard density ρ_n . can be examined and be utilised as further criterion for termination of the iteration.

In the following figures 4 to 10 are outlined now advantageous embodiments of the devices according to the invention according to the claims 12 and 23. Same numerals indicate respectively identical or functionally similar devices, so that in the figures 4 to

SUB
A12
B13

09869782-070501

^{B13}
SUB A10
cont D. 10 In essence only the differences in the respective embodiments are represented and otherwise is referred to the respective above mentioned description.

09869782-070501
5 The photometric device presented in the figure 3a according to the state of art shows a radiation source 2, which emits spectral broadband radiation 15 in the relevant spectral area, here especial in infrared. The radiation 15 is guided through the probe gas 1, that itself is located in a probe cell 3, for example in a optical measuring cell, which is chargeable by means of the orifices 4, 5. The radiation 15 is absorbed wavelength selective by the probe gas 1 corresponding of its composition of substances and the transmitted radiation 16 is fed to a modulation unit 6, that performs a spectral analysis of the transmitted radiation 16. For this, like in the figure 3 schematically hinted, a prism can be used, which discharges the radiation 8 of the different wavelengths in different directions. Also arrangements of gratings or the same are usual. The selective detection of a certain wavelength is achieved by positioning of a radiation sensible radiation receiver 7 along the adjustment direction 14 in the appropriate position of the spectral widened measurement radiation 8 or by way of turning the prism. A computational unit 10 registers the by means of signals 9 transmitted, detected measurement radiation 8 as a function of the wavelength and gains in this way the transmission spectrum according to figure 3b of the probe gas 1, that can be evaluated particularly according to the above presented procedure in respect of the wanted gas quantities. Beside the signal processing the computational unit 10 serves in addition for the control of the radiation source 2 by means of signals 11, for example for modulating of the luminous intensity, as well as for the control of the selection of the wavelength by means of signals 12 according to the movements of the prism 17 or by means of signals 13 according to the movement of the radiation receiver. The device according to figure 3a delivers a transmission spectrum, as for example represented in Fig 3b, that is evaluated according to the above presented procedure in the way of MVA or in the way of DSA and will be translated by means the above presented iterative procedures to the desired standard quantities.

2.14
SUB 211

5 In a first conceivable embodiment of the invention according to claim 12 and figure 4 the device is coupled by means of optical fibres 19 to the probe cell 3 and can like this be installed in an explosion-free room behind an explosion barrier 20, while the intrinsic measuring cell itself is located near the gas conduit 18 within not explosion free room. For the recording of a null spectrum and in doing so the offset-transmission of the optical system the probe cell 3 can be filled with a spectroscopical inactive inert gas 23, in infrared for example nitrogen. Alternatively the probe cell 3 can be admitted in addition with a calibrating gas 24 for the verification of the measurement values. Null- and calibrating measurement are carried out in appropriate selected time intervals and are controlled via a valve block 25 by the computational unit 10. This embodiment can also be combined with the other implementations.

09869782-070501

15 The device according to the invention registers according to figure 4 no spectral continuously resolved spectrum, instead uses for selection of the wavelength optical filter 21 (for example interference filter), which are positioned in front of each radiation receiver 7. Alternative also an individual radiation receiver 7 can be used, whereas the filter 21 is positioned sequentially for example by a wheel of filters 45 into the optical path of a optic 22. The filter regions are for example selected like this, that they emulate the spectral function in the above presented DSA-procedure and deliver directly the wanted quantities in operation condition. The translation to the standard quantities is done by means of the above presented two-stage iterational procedure. An especially easy assembly for the filters would be for example made of four filters: one each for CH₄, higher CH, CO₂ and reference gas.

25 The first embodiment of a device presented in the figure 5 according to the invention according to claim 23 works in combination with a prism- or grating-spectrometer, as well as provided in the embodiment according to the state of art according to figure 3. The selection of the wavelength is done thereby not by means of movements of a prism or of the grating or of the radiation receiver 7, instead by a in a chopper arrangement 28 provided with a sector aperture 46. This sector aperture 46 owns free sectors 30, which

30

Sub
Big
Conc'd

5 sequential set free areas of the outlet slit of the grating 47 or the same and with it selected regions of wavelengths of the spectrum of the measurement radiation 8. A projection lens 22 maps the transmitted measurement radiation 8 to a radiation receiver 7; based on the slit-like geometry for the projection lens 22 for example a cylindrical lens will be the best opportunity. With the information of the position of the rotational direction 29 of the rotating aperture 46 and therewith of the wavelength the computational unit 10 determines a simplified transmission spectrum, which is evaluated with the above presented procedure. By the selection of the position and width of the sectors 30 regions of wavelengths can be selected, which are optimised for the above presented evaluation procedures, particularly for DSA. The computational unit 10 controls with signals 27 also the rotational movement of the aperture 46. An alternative type design of the chopper arrangement 28 according to figure 5 is represented in the figure 6. The chopper arrangement 28 acts appropriately to sector aperture 46 of the figure 5, whereby however not sequential fixed wavelength intervals are faded in, instead the spectrum rather gets scanned continuously by a spiral slit opening 31 arranged in the aperture 46. The advantage is thereby a superior flexibility for the evaluation of the spectrums, however the measurement time for every interval of wavelength is shortened and in doing so the signal-noise is increased. As with the chopper arrangement 28 according to figure 5 the computational unit 10 ascertains based on the position of the aperture 46 the respective present wavelength and in doing so records a transmission spectrum for the above evaluation procedures MVA and DSA.

25 In the figure 7 is represented an enhanced device with a chopper arrangement 28, in which a reference channel is realised by an optical waveguide 35 for that purpose, that the open in the surrounding air guided optical path of the signals by the way of the waveguide 34 and the one of the reference beam through the waveguide 35 are identical, so that for example the underground spectrum of the atmospheric CO₂ as well as variances of the radiation source and the radiation receiver can be ideally
30 suppressed. The change between the two channels in the waveguides 34, 35 is done

thereby by a respective chopper arrangement 28 between radiation source 7 and coupling in of the measurement radiation 8 into the waveguides 34, 35, which illuminates by way of different sector element groups 36, 37 alternating the fibre facets of the two waveguides. The ends of the two channels of the waveguides 34, 35 are
5 arranged closely to each other in the entrance slit of the monochromator 32, so that both fibre facets cover the same wavelength regions and a spreading of the slit and therewith a reduction of the spectral resolution is not required. The figures of the two fibre facets in the leaving slit of the monochromator 32 lie corresponding just as one upon the other and must be unified in the radiation receiver 7; this is done for example
10 by choosing a radiation receiver 7 with a sufficiently large surface, for the sake of capturing both light spots, or by an easily defocussed adjustment of the radiation receiver 7. The intrinsic measuring apparatus can be installed doing this for example in a housing 33 encapsulated against environmental conditions.

15 A further variant of the device according to the invention shows the figure 8. In this embodiment the chopper arrangement 28 simultaneously takes over two functions: on the one hand the selection of the wavelengths by means of segment openings 30 will be carried out corresponding to the embodiments according to the figures 6 and 7, on the other hand the switchover between signal- and reference channel according to figure 7
20 is done by means of respective segment openings 36, 37 in the border area of this same aperture 46. The advantage of this embodiment is the compact and easy carrying out of both functions with only one drive, in addition the synchronousness of the wavelength modulation with the modulation of the beam channel is automatically assured. In the presentation according to the figure 8 in addition a so-called Y-fibre coupler 38 will be
25 used to unify the radiation of reference channel and signal channel in the entrance slit of the monochromator 32.

The figure 9 shows a design of a probe cell 3 in the shape of a hollow shaft guide 39. In optical probe cells 3 with coupling to a waveguide 34 the measurement radiation 8,
30 which is discharged by the waveguide 34, will usually be collimated by an optic and in

the form of a parallel beam guided through the probe cell 3, and is further focussed with an other optic again into the waveguide 34. In Fig 9 the probe cell 3 is realised as optical hollow shaft guide 39. An optical hollow shaft guide 39 is generally a small tube with round cross-section and an inside mirrored wall, so that the coupled-in measurement radiation 8 will be guided through multiple reflection through the hollow shaft guide 39. The dimensioning of the cross-section of the hollow shaft guide 39 is located preferably about in the size range of the waveguide 34. Hereby additional optics for coupling of waveguide 34 and hollow shaft guide 39 will be unnecessary.

The design of the probe cell 3 as hollow shaft guide 39 has thereby following advantages: The probe cell 3 can be coupled as hollow shaft guide 39 very simple and without additional optics to the supplying and the carrying away waveguide 34. The design is essentially more compact as in the case of a free guided parallel beam. The effective path length of the measured length and therefore also the sensitivity by way of the zigzag-path is improved. Because the hollow shaft guide 39 is build as a pipe, he can be connected in addition via connectors 4, 5 very good with gas conduits 18.

A further application of this optical probe cell 3 in form of a hollow shaft guide 39 can be the optical detection for gas chromatographs. The stream of gas, which escapes off the capillary column of the gas chromatograph, is directly guided through the hollow shaft guide 39 and optically determined.

The figure 10b shows a further improvement of the coupling of the waveguide 34 to the probe cell 3 by means of so-called GRIND-lenses 43. In the atmospheric air are contained gases as CO₂ and water steam, which might disturb by their fluctuation the measuring of gas in the probe cell 3 at the state of art according to figure 10a; it has therefore to be avoided, that the measurement radiation 8 is guided across free optical paths 40 with surrounding air. In a spectrometer this effect will be compensated by a two-beam-concept; in the probe cell 3 however as well free paths 40 occur in the fibre optic. By using lenses with a radial gradient of refraction index, so-called GRIND-

lenses 43 according to figure 10b the collimation and focusing can be done without free optical paths 40 and so the influence of the measurement through atmospherical air is suppressed.

5 In the figure 11 is represented an embodiment of the invention with modulation of the measurement radiation, in which respectively for certain spectral regions of the digital signal evaluation designed filters 21, 21' and 21" respective radiation sources 2, 2' and 2" are arranged adjacent and the radiation 15 before passing through the probe gas 1 is unified through an arrangement of beam-splitters 48 for the radiation 15. Doing this the
10 filters 21, 21' and 21" are employed in their filter behaviour in such a way, that they only let pass through spectral regions of the emitted radiation coming from the radiation sources 2, 2' and 2", which are desired for the desired spectral regions for the digital signal evaluation. After the unification of the beams of these three partial beams with the beam-splitter 48 the radiation 15 passes a not nearer described probe cell 3, in
15 which probe gas 1 is included. The escaping measurement radiation 8 after having passed the not further described probe cell 3 will be trapped in already basically known manner by a radiation receiver 7 and reported to the computational unit 10 in also already presented manner. This computational unit 10 controls doing this both the radiation receiver 7 as well as by means of control signals 13 also the radiation sources
20 2, 2' and 2" by means of the control signals 11, whereby both the radiation sources 2, 2' and 2" are variable in such a manner, that a modulation of the radiation sources 2, 2' and 2" in amplitude and/or wavelength can be done. Also it is self evidently thinkable, that an additional modulation unit can be provided, which is here however not farther represented, that modulates the measurement radiation 8 itself after leaving
25 the radiation sources 2, 2' and 2". For stabilization of the measurements according to the figure 11 a further optical path is built up, by means of which partial beams coming off the beam-splitter 48 via mirrors 47 and a beam-splitter 48' shine into a reference detector 49 and there can be utilized for the compensation of unwanted effects. Also the reference detector 49 is connected via control signals 11' with the computational
30 unit 10.

SUB 7
A12
B16

The figures 4 to 11 show in a very simplified representation an equipment basic design of a device according to the claims 12 or 23. In this the description is restricted to the essential courses of the procedures and thereto necessary devices. It can be understood, that the expert can carry out with the teachings according to the invention according to the claims a large number of variations and adaptations, that also are covered by the item of the invention.

09869782-070501

Parts List

	1	-	probe gas
	2, 2', 2''	-	radiation source
	3	-	probe cell
5	4	-	outlet
	5	-	intake
	6	-	modulation unit
	7	-	radiation receiver
	8	-	measurement radiation
10	9	-	measurement signals
	10	-	computational unit
	11	-	control signals for radiation source
	12	-	control signals for modulation unit
	13, 13'	-	control signals for radiation receiver
15	14	-	movement device for radiation receiver
	15	-	radiation in front of probe cell
	16	-	radiation after leaving of probe cell
	17	-	direction of movement of oscillation
	18	-	gas supply
20	19	-	waveguide
	20	-	explosion barrier
	21	-	interference filter
	22	-	projection lens
	23	-	inertial gas
25	24	-	calibrating gas
	25	-	valve block
	26	-	drive for taking probes
	27	-	control signals for chopper
	28	-	chopper

0986978-0705028-28/69860

09869782-070501

- 29 - direction of turning of chopper
- 30 - opening segments of chopper
- 31 - spiral chopper
- 32 - monochromator
- 5 33 - housing
- 34 - first optical waveguide
- 35 - second optical waveguide
- 36 - first sector element group
- 37 - second sector element group
- 10 38 - Y-fibre coupler
- 39 - hollow shaft guide
- 40 - free path
- 41 - parallel beam
- 42 - spherical lens
- 15 43 - GRIND-lens
- 44 - direct coupling
- 45 - filter carrier
- 46 - aperture
- 47 - mirror
- 20 48, 48' - beam-splitter
- 49 - reference detector